

- Application of skin creams
- Debridement of nails
- Other hygienic or preventive maintenance care

### ***Sterilization (ARM 37.86.104)***

#### **Elective Sterilization**

Elective sterilizations are sterilizations done for the purpose of becoming sterile. Medicaid covers elective sterilization for men and women when all of the following requirements are met:

1. Client must complete and sign the *Informed Consent to Sterilization* (MA-38) form at least 30 days, but not more than 180 days, prior to the sterilization procedure. This form is the **only** form Medicaid accepts for elective sterilizations (see *Appendix A Forms* for the form and instructions). If this form is not properly completed, payment will be denied.

The 30-day waiting period may be waived for either of the following reasons:

- **Premature Delivery.** The *Informed Consent to Sterilization* must be completed and signed by the client at least 30 days prior to the estimated delivery date and at least 72 hours prior to the sterilization.
  - **Emergency Abdominal Surgery.** The *Informed Consent to Sterilization* form must be completed and signed by the client at least 72 hours prior to the sterilization procedure.
2. Client must be at least 21 years of age when signing the form.
  3. Client must not have been declared *mentally incompetent* (see *Definitions*) by a federal, state or local court, unless the client has been declared competent to specifically consent to sterilization.
  4. Client must not be confined under civil or criminal status in a correctional or rehabilitative facility, including a psychiatric hospital or other correctional facility for the treatment of the mentally ill.

Before performing a sterilization, the following requirements must be met:

- The client must have the opportunity to have questions regarding the sterilization procedure answered to his/her satisfaction.
- The client must be informed of his/her right to withdraw or withhold consent anytime before the sterilization without being subject to retribution or loss of benefits.
- The client must be made aware of available alternatives of birth control and family planning.



All forms required for sterilizations can be copied from *Appendix A Forms*, can be ordered using the *Medicaid Form Order* sheet in the *General Information For Providers* manual, or can be downloaded from the Provider Information Web Site (see *Key Contacts*.)



Medicaid covers hysterectomies only when they are a result of a procedure performed to address another medical problem, not when the primary purpose is to render the client sterile.

- The client must understand the sterilization procedure being considered is irreversible.
- The client must be made aware of the discomforts and risks which may accompany the sterilization procedure being considered.
- The client must be informed of the benefits and advantages of the sterilization procedure.
- The client must know that he/she must have at least 30 days to reconsider his/her decision to be sterilized.
- An interpreter must be present and sign for those clients who are blind, deaf, or do not understand the language to assure the person has been informed.

Informed consent for sterilization may not be obtained under the following circumstances:

- If the client is in labor or childbirth.
- If the client is seeking or obtaining an abortion.
- If the client is under the influence of alcohol or other substance which affects his/her awareness.

### Medically Necessary Sterilization

When sterilization results from a procedure performed to address another medical problem, it is considered a medically necessary sterilization. These procedures include hysterectomies, oophorectomies, salpingectomies and ochiectomies. Every claim submitted to Medicaid for a medically necessary sterilization must be accompanied by one of the following:

- A completed *Medicaid Hysterectomy Acknowledgement* form (MA- 39) for each provider submitting a claim. See *Appendix A Forms* for the form and instructions. It is the provider's responsibility to obtain a copy of the form from the primary or attending physician. The client must sign and date this form at least 30 days prior to the procedure (see 42 CFR 441.250 for the federal policy on hysterectomies and sterilizations).
- For clients who have become retroactively eligible for Medicaid, the physician must certify in writing that the surgery was performed for medical reasons and must document one of the following:
  - The individual was informed prior to the hysterectomy that the operation would render the client permanently incapable of reproducing.
  - The reason for the hysterectomy was a life-threatening emergency.
  - The client was already sterile at the time of the hysterectomy and the reason for prior sterility.

A notation "Not a Sterilization" on a claim is not sufficient to fulfill these certification requirements.

### PA Criteria for Specific Services (continued)

Service	PA Contact	Documentation Requirements										
• Reduction Mammo-plasty	<p>SURS P.O. Box 202953 Helena, MT 59620-2953</p> <p><b>Phone:</b> (406) 444-0190 Helena and out of state</p> <p>(406) 444-1441 Helena and out of state</p> <p><b>Fax:</b> (406) 444-0778</p>	<ul style="list-style-type: none"><li>Both the Referring physician and the surgeon must submit documentation.</li><li>Back pain must have been documented and present for at least six months, and causes other than weight of breasts must have been excluded.</li><li><b>Indications for female client:</b></li><li>Contraindicated for pregnant women and lactating mothers. A client must wait six months after the cessation of breast feeding before requesting this procedure.</li><li>Female client 16 years or older with a body weight less than 1.2 times the ideal weight.</li><li>There must be severe, documented secondary effects of large breasts, unresponsive to standard medical therapy administered over at least a six month period. This must include at least two of the following conditions:<ul style="list-style-type: none"><li>Upper back, neck, shoulder pain that has been unresponsive to at least six months of documented and supervised physical therapy and strengthening exercises</li><li>Paresthesia radiating into the arms. If parathesia is present, a nerve conduction study must be submitted.</li><li>Chronic intertrigo (a superficial dermatitis) unresponsive to conservative measures such as absorbent material or topical antibiotic therapy. Document extent and duration of dermatological conditions requiring antimicrobial therapy.</li><li>Significant shoulder grooving unresponsive to conservative management with proper use of appropriate foundation garments which spread the tension of the support and lift function evenly over the shoulder, neck and upper back.</li></ul></li></ul> <p>Documentation in the client’s record must indicate and support the following:</p> <ul style="list-style-type: none"><li>History of the client’s symptoms related to large, pendulous breasts.</li><li>The duration of the symptoms of at least six months and the lack of success of other therapeutic measures (e.g., documented weight loss programs with six months of food and calorie intake diary, medications for back/neck pain, etc.).</li><li>Guidelines for the anticipated weight of breast tissue removed from each breast related to the client’s height (which must be documented):</li></ul> <table><tr><th>Height</th><th>Weight of tissue per breast</th></tr><tr><td>less than 5 feet</td><td>250 grams</td></tr><tr><td>5 feet to 5 feet, 2 inches</td><td>350 grams</td></tr><tr><td>5 feet, 2 inches to 5 feet, 4 inches</td><td>450 grams</td></tr><tr><td>greater than 5 feet, 4 inches</td><td>500 grams</td></tr></table> <ul style="list-style-type: none"><li>Pre-operative photographs of the pectoral girdle showing changes related to macro-mastia.</li><li>Medication use history. Breast enlargements may be caused by various medications (e.g., sironolactone, cimetidine) or illicit drug abuse (e.g., marijuana, heroin, steroids). Although rare in women, drug effects should be considered as causes of breast enlargement prior to surgical treatment since the problem may recur after the surgery if the drugs are continued. Increased prolactin levels can cause breast enlargement (rare). Liver disease, adrenal or pituitary tumors may also cause breast enlargement and should also be considered prior to surgery.</li><li><b>Indications for male client:</b></li><li>If the condition persists, a client may be considered a good candidate for surgery. Clients who are alcoholic, illicit drug abusers (e.g., steroids, heroin, marijuana) or overweight are not good candidates for the reduction procedure until they attempt to correct their medical problem first.</li><li>Documentation required: length of time gynecomastia has been present, height, weight, and age of the client, pre-operative photographs</li></ul>	Height	Weight of tissue per breast	less than 5 feet	250 grams	5 feet to 5 feet, 2 inches	350 grams	5 feet, 2 inches to 5 feet, 4 inches	450 grams	greater than 5 feet, 4 inches	500 grams
Height	Weight of tissue per breast											
less than 5 feet	250 grams											
5 feet to 5 feet, 2 inches	350 grams											
5 feet, 2 inches to 5 feet, 4 inches	450 grams											
greater than 5 feet, 4 inches	500 grams											

<b>PA Criteria for Prescription Drugs</b>	
<b>Drug</b>	<b>Criteria</b>
Non-steroidal Anti-Inflammatory Drugs  PA required for all single-source NSAIDS: Ponstel Mobic Naprelan	Trial and failure with at least <u>two</u> multiple-source products must be documented.
Celebrex (celecoxib) Vioxx (rofecoxib) Bextra (valdecoxib)	No history of aspirin sensitivity or allergy to aspirin or other NSAID, and/or aspirin triad, and at least one of the following: <ul style="list-style-type: none"> <li>• History of previous GI bleeding within the last 5 years</li> <li>• Current or recurrent gastric ulceration</li> <li>• History of NSAID-induced gastropathy</li> <li>• Currently treated for GERD</li> </ul> For clients 65 and older, no concomitant aspirin use. Vioxx 50mg is not recommended for chronic use. Medicaid does not cover Vioxx at this dose for extended periods.
Disease-modifying anti-rheumatic drugs (DMARD) Arava (leflunomide) Enbrel (etanercept) Humira (adalimumab) Kineret (anakinra) Remicade (infliximab)	<ul style="list-style-type: none"> <li>• Diagnosis of rheumatoid arthritis</li> <li>• Rheumatology consult with date and copy of consult included</li> <li>• Failure with or contraindication to methotrexate alone or in combination with sulfasalazine, hydroxychloroquine or Arava, in which case Enbrel, Remicade, or Kineret may be approved either alone or in combination with Arava.</li> <li>• Kineret may be used alone or in combination with DMARD's other than tumor necrosis factor (TNF) blocking agents (i.e. Enbrel)               <ul style="list-style-type: none"> <li>• Enbrel or Remicade whether alone or in combination with methotrexate or Arava may be approved for first-line treatment in patients with moderately to severely active rheumatoid arthritis as evidenced by:                   <ul style="list-style-type: none"> <li>• &gt; 10 swollen joints</li> <li>• <math>\geq</math> 12 tender joints</li> <li>• Elevated serum rheumatoid factor levels or erosions on baseline x-rays</li> </ul> </li> </ul> </li> </ul>
Remicade (infliximab)	Diagnosis of: <ul style="list-style-type: none"> <li>• Moderately to severely active Crohn's disease for patients with an inadequate response to conventional therapy</li> <li>• Fistulizing Crohn's disease</li> </ul>
Ambien (zolpidem) Sonata (zaleplon) Quantity limited to 15 tablets per month.	Trial and failure with at least <u>two</u> multi-source prescription sleep-inducing drugs.
Oxycodone HCL Controlled-Release (OxyContin)	<ul style="list-style-type: none"> <li>• Diagnosis of oncologic pain</li> <li>• Prior authorization is required for all dosing above twice a day and above 320 mg per day.</li> </ul>

<b>PA Criteria for Prescription Drugs (continued)</b>	
<b>Drug</b>	<b>Criteria</b>
Thalomid (thalomide)	Treatment of the cutaneous manifestations of moderate-to-severe erythema nodosum leprosum (ENL). Considered for other diagnoses on individual basis.
Zyvox (linezolid)	Adult patients with vancomycin-resistant enterococcus.
Zoloft 25 mg & 50 mg (sertraline)	Authorized for patients requiring dosages where tab splitting would be inappropriate (i.e., 75 mg, 125 mg).
Tretinoin PA required for patients 26 years and older.	Diagnose of: <ul style="list-style-type: none"> <li>• Skin cancer</li> <li>• Lamellar ichthyosis</li> <li>• Darier-White disease</li> <li>• Psoriasis</li> <li>• Severe recalcitrant (nodulocystic) acne</li> </ul>
Growth hormones	<p>Diagnosis of:</p> <ul style="list-style-type: none"> <li>• Growth hormone deficiency in children and adults</li> <li>• Growth retardation of chronic renal insufficiency</li> <li>• Turner's Syndrome</li> <li>• AIDS-related wasting</li> </ul> <p>Children and adolescents must meet the following criteria:</p> <ul style="list-style-type: none"> <li>• Standard deviation of 2.0 or more below mean height for chronological age</li> <li>• No expanding intracranial lesion or tumor diagnosed by MRI</li> <li>• Growth rate below five centimeters per year</li> <li>• Bone age 14-15 years or less in females and 15-16 years or less in males</li> <li>• Epiphyses open</li> </ul> <p><b>Growth hormone deficiency in children:</b> Failure of any two stimuli tests to raise the serum growth hormone level above 10 nanograms/milliliter.</p> <p><b>Growth retardation of chronic renal insufficiency:</b> Irreversible renal insufficiency with a creatinine clearance &lt;75 ml/min per 1.73m<sup>2</sup> but pre-renal transplant.</p> <p><b>Turner's Syndrome:</b> Chromosomal abnormality showing Turner's syndrome.</p> <p><b>Growth hormone deficiency in adults:</b></p> <ul style="list-style-type: none"> <li>• <b>Adult Onset:</b> Patients have somatotropin deficiency syndrome (SDS) either alone or with multiple hormone deficiencies, (hypopituitarism), as a result of pituitary disease, hypothalamic disease, surgery, radiation therapy or trauma.</li> <li>• <b>Childhood Onset:</b> Patients who had growth hormone deficient during childhood and now have somatotropin deficiency syndrome (SDS).</li> </ul>

<b>PA Criteria for Prescription Drugs (continued)</b>	
<b>Drug</b>	<b>Criteria</b>
Dipyridamole	As adjunct to warfarin anticoagulants in the prevention of postoperative thromboembolic complications of cardiac valve replacement.
<p>Gastro-intestinal drugs</p> <p>Includes H-2 antagonists, proton pump inhibitors, Sucralfate, and Cytotec</p> <p>PA required after a 90-consecutive-day course of treatment with proton pump inhibitors.</p> <p>PA required after 30 consecutive days with Sucralfate.</p> <p>Concurrent therapy requires PA.</p> <p>Consecutive alternating regimens of different drugs counted as part of the total 90-day period.</p>	<p>Diagnosis of:</p> <ul style="list-style-type: none"> <li>• Hypersecretory conditions (Zollinger-Ellison syndrome, systemic mastocytosis, multiple endocrine adenomas)</li> <li>• Symptomatic gastroesophageal reflux (not responding or failure of maintenance therapy)</li> <li>• Symptomatic relapses (duodenal or gastric ulcer) on maintenance therapy</li> <li>• Barretts esophagus</li> <li>• GERD</li> </ul> <p>Other conditions considered on an individual basis.</p> <p>Sucralfate authorized for an initial 30-day concurrent therapy with histamine H2-receptor antagonists and proton pump inhibitors. Concurrent therapy for a period exceeding 30 days considered duplicate therapy and not covered.</p> <p>Concurrent, combination therapy of proton pump inhibitors and histamine H2-receptor antagonists regarded as duplicate therapy and not covered.</p>
<p>Migraine Headache Drugs</p> <p>For monthly quantities greater than 9 tablets:</p> <p>Imitrex (sumatriptan): 4 injections (2 kits) <b>or</b> 6 units of nasal spray</p> <p>Maxalt (rizatriptan)</p> <p>Zomig (zolmitriptan) and Zomig ZMT (zolmitriptan)</p> <p>Migranal (dihydroergotamine): 4 units</p> <p>Axert (almotriptan)</p> <p>Frova (frovatriptan)</p> <p>Relpax (electriptan)</p> <p>Amerge (naratriptan HCl)</p>	<p>Indicated only for treatment of <u>acute</u>, migraine/cluster headache attacks for patients who meet the following criteria:</p> <ul style="list-style-type: none"> <li>• No history of, or signs or symptoms consistent with, ischemic heart disease (angina pectoris, history of MI or documented silent ischemia) or Prinzmetal's angina</li> <li>• No uncontrolled hypertension</li> <li>• No complicated migraine including vertebrobasilar migraine</li> <li>• Not pregnant</li> <li>• No use of ergotamine-containing medication(s) within previous 24-hours</li> <li>• No use of MAOI within previous 2-weeks</li> <li>• Non-responsive to NSAIDS, acetaminophen, combination analgesics (isometheptene, butalbital, +/- metoclopramide), or ergot derivatives, or these medications are contraindicated</li> </ul> <p>Only one migraine medication may be prescribed within a month.</p> <p>Concurrent therapy with Stadol will not be covered.</p>

<b>PA Criteria for Prescription Drugs (continued)</b>	
<b>Drug</b>	<b>Criteria</b>
Aggrenox (aspirin/dipyridamole)	For prevention of recurrent stroke in patients who have experienced a transient ischemic attack or previous ischemic stroke and who have had a recurrent stroke while on aspirin or have failed plavix and have a contraindication to aspirin.
Toradolral (ketorolac) For quantity greater than a 5-day supply within a month	Indicated for the short-term treatment of acute pain. Authorization considered on an individual basis.
Stadol (butorphanol) PA required for quantities greater than 3 - 2.5 ml metered dose spray pumps within a one-month period	Indicated for management of pain including post-operative analgesia or acute migraine headache pain for patients who meet the following criteria: <ul style="list-style-type: none"> <li>• No history of hypersensitivity to butorphanol or any components of the product</li> <li>• No history of narcotic dependency or abuse</li> <li>• Not pregnant</li> <li>• No ulcerations of the nasal mucosa</li> <li>• No history of psychological or neurological disorder</li> <li>• No history of head trauma within the previous 7 days</li> <li>• 18 years of age or older</li> <li>• Non-responsive to NSAIDS, acetaminophen, combination analgesics (isometheptene, butalbital, +/- metoclopramide), or ergot derivatives, or these medications are contraindicated.</li> </ul>
Smoking Cessation Drugs  Nicotine-replacement products Zyban (bupropion)	Authorization given for 4-month course of therapy. Four trials of therapy are allowed.
Trental (pentoxifylline) Pletal (cilostazol)  For greater than 12-week supply within a 12-month period.	<ul style="list-style-type: none"> <li>• Diagnosis of <u>intermittent claudication</u> as the result of chronic occlusive arterial disease (COAD) of the lower limbs. Possible causes of COAD include: arteriosclerosis obliterans, thromboangiitis obliterans (Buerger's disease), arteritis, trauma, congenital arterial narrowing, or other forms of peripheral vascular disease resulting in chronic vascular occlusion in the legs; <b>and</b></li> <li>• The patient has shown clinical improvement in their COAD while on pentoxifylline or cilostazol.</li> <li>• Considered on an individual basis when pentoxifylline or cilostazol is being used as part of a standardized treatment protocol, e.g. bone marrow or oncology treatment protocols.</li> </ul>
Viagra (sildenafil) Quantity limited to one (1) tablet per month	<ul style="list-style-type: none"> <li>• Diagnosis of erectile dysfunction.</li> <li>• Males only, 18 years of age or older.</li> <li>• No concomitant organic nitrate therapy.</li> </ul>

<b>PA Criteria for MHSP Prescription Drugs</b>	
<b>Drug</b>	<b>Criteria</b>
buspirone (Buspar)	<ul style="list-style-type: none"> <li>• Augmentation of depression and/or obsessive compulsive disorder (OCD).</li> <li>• Generalized anxiety disorder.</li> </ul>
zaleplon (Sonata) zolpidem (Ambien)	Trial and failure with at least <b>two</b> multi-source prescription sleep-inducing drugs.
amotrigine (Lamictal)	<ul style="list-style-type: none"> <li>• Diagnosis of bi-polar disorder.</li> <li>• Intolerance, contraindication, or partial response to Lithium, Tegretol, or Depakote.</li> </ul>
guanfacine (Tenex) isradipine (DynaCirc) levothyroxine sodium (Synthroid) liothyronine sodium (Cytomel) pindolol (Visken) propranolol HCl (Inderal) verapamil, verapamil HCl (Calan)	Use as augmentation strategy specifically related to mental health treatment.
maprotiline HCl (Ludiomil)	Considered on an individual basis.
sertraline (Zoloft 25 mg & 50 mg )	Authorized for patients requiring dosages where tablet splitting would be inappropriate (i.e., 75 mg, 125 mg).
gabapentin (Neurontin)	Must specify if anxiety (generalized anxiety, panic disorder, post traumatic stress disorder) and/or compelling reason with bipolar disorder.
topiramate (Topamax)	Diagnosis of bipolar disorder, obesity, intolerance, time effective of Lithium, Depakote, Trileptal/Tegretol.



- For services where codes or definitions differ between the CPT-4 and the *American Society of Anesthesiologists' Relative Value Guide*, Medicaid adopts the CPT-4 version.
- Include the total number of minutes in field 24g (*Days or Units*) on the CMS-1500 claim form. Medicaid will convert the number of minutes to the number of time units. Do not include the base units on the CMS-1500 as the claims processing system determines the number of base units (see the *Completing a Claim* chapter in this manual).

### ***Bundled services***

Certain services with CPT-4 codes (eg., telephone advice, some pulse oximetry services) are covered by Medicaid but have a fee of zero. This means that the service is typically “bundled” with an office visit or other service. Since the bundled service is covered by Medicaid, providers may not bill the client separately for it.

### ***Cosmetic services***

Include the prior authorization number in field 23 (*Prior Authorization Number*) on the CMS-1500 claim form (see the *Completing a Claim* chapter in this manual).

### ***EPSDT Well Child Screens***

- Bill for a complete screen using the appropriate evaluation and management (E&M) code for preventive medicine services.
- When billing for partial screens, use the appropriate preventive medicine code with modifier 52 (reduced services).
- See also the EPSDT chapter in this manual.
- For EPSDT overrides on limits and cost sharing, see the *Completing a Claim* chapter in this manual.

### ***Family planning services***

Contraceptive supplies and reproductive health items provided free to family planning clinics cannot be billed to Medicaid. When these supplies are not free to the clinic, providers associated with a family planning clinic can bill Medicaid for these items using local code Z0695. These items include:

- |                    |                        |
|--------------------|------------------------|
| • Diaphragm        | • Oral contraceptive   |
| • Foam jelly/cream | • Terazol              |
| • Condoms          | • Bichloracetic acid   |
| • Monistat         | • Trichloroacetic acid |
| • Sultrin          | • Contraceptive film   |

For family planning overrides on cost sharing, see the *Completing a Claim* chapter in this manual.

### ***Immunizations***

- Use code Z0805 to bill for the administration of vaccines under the Vaccines for Children (VFC) program.
- There must be a VFC covered code for each unit of service billed with code Z0805. For a list of VFC covered vaccines, contact the Department's immunization program at (406) 444-5580.
- No more than four diagnosis codes are necessary.
- Bill each VFC code with \$0.00 charges.

For example, a provider administers two vaccines: MMR and pneumococcal conjugate. The provider's charge for each vaccine administration is \$5.00.

24.	A	B	C	D	E	F	G	H	I	J	K
	DATE(S) OF SERVICE	Place of Service	Type of Service	PROCEDURES, SERVICES, OR SUPPLIES (Explain Unusual Circumstances)	DIAGNOSIS CODE	\$ CHARGES	DAYS OR UNITS	EP/SDT Family Plan	EMG	COB	RESERVED FOR LOCAL USE
	MM DD YY	MM DD YY		CPT/HCPCS MODIFIER							
1	05   13   02		11 0	Z0805	1	10.00	2				
2	05   13   02		11 0	90707	1	0.00	1				
3	05   13   02		11 0	90669	1	0.00	1				

### ***Obstetrical services***

If the provider's care includes prenatal (antepartum) and/or postnatal (postpartum) care in addition to the delivery, the appropriate global OB code must be billed. Antepartum care includes all visits until delivery, and there are different codes for specified numbers of visits. There are also different codes for antepartum and postpartum care when only one or the other is provided. Please review your CPT coding book carefully.

### ***Reference lab billing***

Under federal regulations, all lab services must be billed to Medicaid by the lab that performed the service. Modifier 90, used to indicate reference lab services, is not covered by Medicaid.

### ***Sterilization***

- For elective sterilizations, a completed *Informed Consent to Sterilization* (MA-38) form must be attached to the claim for each provider involved or payment will be denied. This form must be legible, complete, and accurate, and revisions are not accepted. It is the provider's responsibility to obtain a copy of the form from the primary or attending physician.
- For medically necessary sterilizations (including hysterectomies), one of the following must be attached to the claim, or payment will be denied:
  - A completed *Medicaid Hysterectomy Acknowledgement* form (MA- 39) for each provider submitting a claim. See *Appendix A Forms* for the form and instructions. It is the provider's responsibility to obtain a copy of the form from the primary or attending physician. The client must sign and date this form at least 30 days prior to the procedure (see 42 CFR 441.250 for the federal policy on hysterectomies and sterilizations).